

PROCESS FOR TREATING FLUE GAS**Publication number:** EP1089804**Publication date:** 2001-04-11**Inventor:** LAGANA VINCENZO (IT)**Applicant:** SIIRTEC NIGI S P A (IT)**Classification:****- International:** *B01D53/56; C01C1/02; C01C1/08; B01D53/56; C01C1/00; (IPC1-7): B01D53/56***- European:** B01D53/56; C01C1/02; C01C1/08D**Application number:** EP19990921144 19990514**Priority number(s):** WO1999IT00135 19990514; IT1998MI01155 19980525**Also published as:**

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An improved process for combustion flue gas conditioning in which ammonia is injected into a stream of said combustion flue gas, wherein urea is dissolved in water, forming a dissolved urea solution. The dissolved urea solution is heated and pressurized and the dissolved urea therein hydrolyzed, forming ammonia and carbon dioxide. The ammonia and carbon dioxide are then stripped from the resulting hydrolysis solution and injected into the stream of combustion flue gas to be treated. Water from the stripped hydrolysis solution is then recycled for use in dissolving fresh urea.

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<p>(21) International Application Number: PCT/IT99/00135</p> <p>(22) International Filing Date: 14 May 1999 (14.05.99)</p> <p>(30) Priority Data: MI98A001155 25 May 1998 (25.05.98) IT</p> <p>(71) Applicant (for all designated States except US): SI-IRTEC-NIGI S.P.A. [IT/IT]; Via Algardi, 2, I-20148 Milano (IT).</p> <p>(72) Inventor; and (75) Inventor/Applicant (for US only): LAGANA', Vincenzo [IT/IT]; Via Sismondi, 43, I-20133 Milano (IT).</p> <p>(74) Agent: CIONI, Carlo; Studio Cioni & Pipparelli, Viale Caldara, 38, I-20122 Milano (IT).</p>		<p>(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).</p> <p>Published <i>With international search report.</i></p>
<p>(54) Title: PROCESS FOR TREATING FLUE GAS</p> <div data-bbox="337 1140 1239 1728"> </div> <p>(57) Abstract</p> <p>Process for combustion flue gas conditioning, by injecting ammonia, in a very simple and viable way, in a stream of said combustion flue gas stream, characterized in that dissolved urea is hydrolyzed by heating under pressure to yield ammonia and carbon dioxide the hydrolysis product are stripped from the hydrolyzed solution by means of hot steam and injected into the flue gas stream while water coming out of the hydrolysed solution is reused to dissolve fresh urea to be fed to the plant.</p>		

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PROCESS FOR TREATING FLUE GAS

Field of the invention

This invention refers to a process for treating flue gases and, more specifically, to a process for purifying (conditioning) such flue gases by ammonia addition. More particularly, this invention relates to a process for flue gas treatment by ammonia addition, whereby the ammonia is derived from hydrolysis of urea.

Description of prior art

It is well known to add ammonia to flue gases to remove, for instance, nitrogen oxides, which are formed in the course of combustion itself. Reaction between ammonia and nitrogen oxides, in the presence of oxygen, results in nitrogen and water formation in accordance with the following reaction:



Ammonia gas is injected into the flue gas stream to be treated by vapor obtained by stripping of the ammonia contained in an ammonia aqueous solution in which the ammonia concentration is roughly 30% by weight. The ammonia aqueous solution has to be transported from the production plant to the user plant, stored, sent to stripping plant to yield the desired ammonia, then the ammonia containing residual water has to be neutralized before its disposal.

Ammonia solutions are presently classified as toxic materials. Consequently, certain specific rules have to be followed for its handling, due to the high volatility and pollution level of ammonia.

In order to avoid the aforementioned problems, use of non-toxic, harmless compounds has been proposed which, under the reaction conditions, may yield ammonia. For this purpose, urea has been used to generate ammonia without the formation of any undesired by-product. Use of urea as conditioning agent of flue

gases solved the problem connected with highly toxic material handling and storage; however, to obtain fine particles of solid urea and uniformly inject it into a flue gas stream still presents substantial problems.

In the US-520688 to Von Harpe, it has been also proposed to employ, as a source of ammonia, instead of solid urea, the products derived from the hydrolysis of urea. Such hydrolysis products are obtained by heating a urea solution. The overall hydrolysis products, namely ammonium carbonate, ammonium carbamate, ammonium bicarbonate and ammonia are used for NO_x reduction. However the simple heating of a concentrated urea solution does not allow to reach a satisfactory amount of hydrolysis products. The above Patent disclosed the use of a basic or acidic environment or addition of a catalyst. However the use of catalyst or acidic or basic products may lead to some problems due to the presence of said material both in the hydrolysis section and NO_x reduction section.

According to US-A-4168299 removal of urea hydrolysis products can be obtained by adsorbing pressurized CO₂ in a diluted urea solution then stripping-off the absorbed CO₂. This process employs a diluted urea solution and a source of a pressurized CO₂.

Summary of the invention

It is one object of the present invention to provide a process for flue gas conditioning by injection of ammonia, in a very simple and viable way, into a stream of said flue gases which avoids problems associated with hazardous material handling and storage.

It is a further object of the present invention to provide a process for *in-situ* production of gaseous ammonia, wherein said ammonia can be used without being subject to any particular safety rules and without any obligation to maintain, transport

and store large volumes of ammonia solutions and toxic noxious material as required by conventional processes, nor to dispose any waste ammonia or ammonia precursor solution.

These and other objects of the present invention are achieved by a process for flue gas conditioning in which urea is dissolved in water to obtain a concentrated urea solution and the dissolved urea is hydrolyzed by heating the concentrated urea solution under pressure, yielding hydrolysis products of ammonia and carbon dioxide. The hydrolysis products are stripped from the solution using hot steam and injected into the flue gas stream while hydrolysis solution coming out of the stripping section, still containing some urea is reused in addition to fresh water, to dissolve fresh urea to be fed to the plant.

Urea solution to be hydrolyzed contains in the range of about 10% to 70% urea by weight. The urea hydrolysis is preferably carried out at a temperature in the range of about 100 to 233°C and a pressure in the range of about 0.1×10^3 KPa to 3.0×10^3 KPa. The hydrolysis solution, after stripping of the gaseous hydrolysis products still contains about 1% to 5% by weight of unhydrolyzed urea.

Brief description of the drawings

These and other objects and features of this invention will be better understood from the following detailed description taken in conjunction with the drawings wherein:

Fig. 1 is a schematic diagram of the process in accordance with one embodiment of this invention.

Description of preferred embodiments

This invention will now be described in more details with reference to Fig. 1 and the following experimental example. The drawing and example are provided to better illustrate the invention and are not to be construed as limiting of the scope thereof.

As shown in Fig. 1, solid urea, in form of prills or crystals, is introduced into hopper 1 having a sufficient capacity for at least one day of continuous hydrolyze operations. Urea, through roll-type feeder 3, is introduced into dissolve 4 where a urea solution of a concentration of about 40% is formed using water condensate from plant battery limits and/or exhausted condensate recycle, which is also introduced into dissolver 4. Urea dissolving operations are facilitated by the action of stirrer 5. Slide valve 2 may be provided at the outlet of hopper 1, if necessary.

The urea/water solution thus obtained is compressed by pump 6 up to hydrolyzer operating pressure and preheated to hydrolysis temperature by means of preheater 7.

10 In accordance with one embodiment of this invention, hydrolyzer operating conditions are as follows:

Pressure	1.9×10^3 KPa
Temperature	195°C
Residence time	40 minutes

15 Under these conditions, about 30%-40% by weight urea concentration in the feedstock decreases to about 1%-5% by weight in the hydrolyzed solution. Ammonia generated during the hydrolysis process is stripped out from the water solution by means of a defined amount of steam coming from battery limits and injected into bottom of hydrolyzer 8. Hydrolysis products comprising water-saturated ammonia and carbon dioxide are conveyed to reaction zone through line 10. Water is expanded through valve 11 and sent to separator 9, operating at atmospheric pressure, and then recycled to the dissolver 4. Vapors derived mainly from ammonia and water expansion through line 12 are added to the hydrolysis vapors in order to avoid any environmental pollution. Water from separator 9 is recycled through line 13 to the

20

25 dissolver 4 at a temperature of about 100°C so that, after having supplied urea with

the heat of solution, a solution is obtained at a temperature of about 40°C.

Example

To better illustrate the process of this invention, a quantitative example is given hereinafter based on an ammonia capacity of 100kg/h.

- 5 The amount of urea fed to the plant is 177 kg/h and the condensate amount coming from plant battery limits is 87kg/h. The water recycle amounts is 150kg/h including about 4.5kg/h of residual urea in the solution. The solution coming out of the dissolver 4 is at a temperature of about 40°C and a urea concentration of about 44% by weight is sent to pump 6, increasing the pressure up to about 1.6×10^3 KPa and,
- 10 through preheater 7, increasing the temperature of about 195°C. The solution enters hydrolyzer 8 resulting in the formation of hydrolysis products having the following characteristics:

Temperature	195°C		
Pressure	1.5×10^3 KPa		
15 Composition	NH ₃	100 kg/h	24.12% by weight
	CO ₂	129 kg/h	31.24% by weight
	<u>H₂O</u>	<u>185 kg/h</u>	<u>44.64% by weight</u>
	Total	414 kg/h	100.00

- Direct steam flow to hydrolyzer 8 required to maintain a constant hydrolysis
- 20 temperature and to strip reaction products is about 155kg/h.

- The result of the process of this invention is that the plant requires only storage facilities for urea, the most used nitrogen fertilizer, which is not subject to any strict regulation and therefore it does not require any special handling care. Other major advantages of the present invention are: safety plant operations, absence of any
- 25 storage by operating personnel of dangerous substances, absence of any polluting

material, and investment cost reduction compared with the use of ammonia solutions according to the prior art.

CLAIMS

1. In a process for combustion flue gas conditioning in which ammonia is injected into a stream of said combustion flue gas stream, the improvement comprising:
dissolving urea in water, forming a dissolved urea solution;
5 heating and pressurizing said dissolved urea solution, forming a heated and pressurized dissolved urea solution;
hydrolyzing the dissolved urea in said heated and pressurized dissolved urea solution, forming a hydrolysed solution comprising ammonia and carbon dioxide;
stripping said ammonia and carbon dioxide from said hydrolysed solution by
10 contacting said hydrolysed solution with steam, leaving behind a stripped hydrolysed solution;
injecting said ammonia and carbon dioxide into said stream of said combustion flue gas; and
recycling said stripped hydrolysed solution for dissolving said urea.
- 15 2. A process in accordance with claim 1, wherein said dissolved urea solution comprises in a range of about 10 to about 70% by weight urea.
3. A process in accordance with claim 1, wherein said dissolved urea is hydrolyzed at a temperature in a temperature range of about 100°C to about 233°C.
4. A process in accordance with claim 1, wherein said dissolved urea is hydrolyzed at
20 a pressure in a pressure range of about 0.1×10^3 KPa to about 3.0×10^3 KPa.
5. A process in accordance with claim 1, wherein said stripped hydrolysed solution is expanded, forming hydrolysis vapors which are combined with said ammonia and carbon dioxide and injected into said stream of said combustion flue gas.
6. A process in accordance with claim 2, wherein said dissolved urea is hydrolyzed at
25 a temperature in a temperature range of about 100°C to about 233°C.

7. A process in accordance with claim 6, wherein said dissolved urea is hydrolyzed at a pressure in a pressure range of about $0,1 \times 10^3$ KPa to about 1.0×10^3 KPa.
8. A process in accordance with claim 7, wherein said stripped hydrolysed solution is expanded, forming hydrolysis vapors which are combined with said ammonia and
- 5 carbon dioxide and injected into said stream of said combustion flue gas.

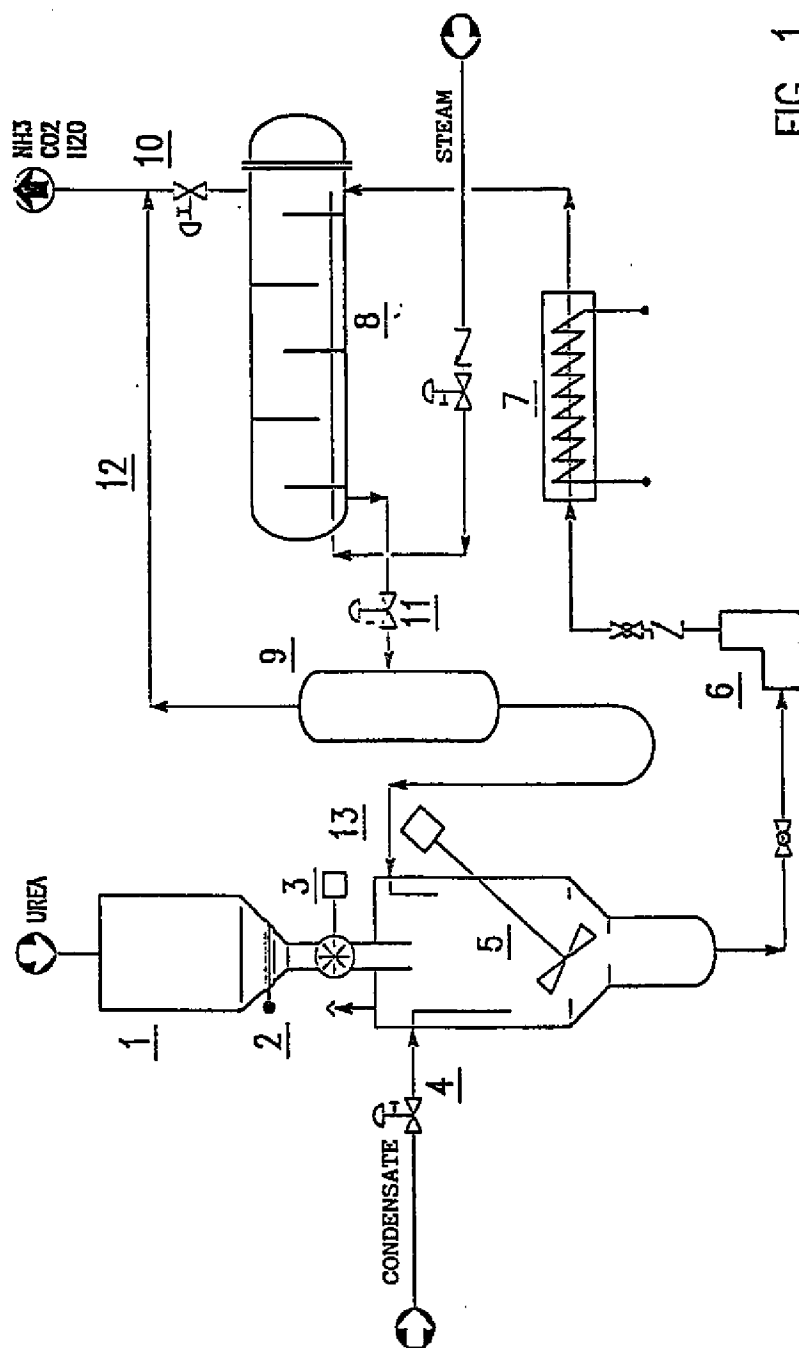


FIG. 1

INTERNATIONAL SEARCH REPORT

International Application No
PCT/IT 99/00135

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 B01D53/56

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 B01D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, X	WO 98 42623 A (COOPER ET AL) 1 October 1998 (1998-10-01) page 20, line 14 - page 27, line 6; examples 1-5	1-8
A	WO 94 13391 A (NALCO FUEL TECH) 23 June 1994 (1994-06-23) page 6, line 5 - page 27, line 26	1-8
A	WO 92 02291 A (FUEL TECH GMBH) 20 February 1992 (1992-02-20) cited in the application page 5, line 28 - page 8, line 19 page 12, line 1 - page 13, line 13; example 1	1-3, 5, 6, 8

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☒ Patent family members are listed in annex.

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30 August 1999

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Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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Information on patent family members

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